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03.10.78-JA-121801 (09.04.80) A61k-09/22
Preparing slew release pharmoceutical - by coating granules contg. oative component with wax, water-sol, high mol. cpd. and nonionic etc. EXAMPLE None given. (7ppW5). surfactant and tabletting

Prepn. of slow release medicine comprises coating instant-Prepn. of slow release medicine comprises coating instant-neously disintegrating granules, which contain a major amt. of medicinal ingredient, with a film composed of wax, water-soluble high molecular compound and nonionic infractant of HLE below 9, and tabletting the coated granules.

Granules which normally release the medicine in 1-10 USE/ADVANTAGE minutes can be treated to release the medicine very slowly over 7-24 hours. Further the releasing velocity of the medicine is independent of pH and is almost constant.

DETAILS

Usually a film of thickness 15-50 p is coated on the granules. The water-soluble high molecular compound (the water solubility of which is independent of pli) can be (the water solubility of which is independent of pli) can be the water solubility of which is independent of pli) can be the water soluble that the water soluble independent of pli) can be the water soluble that the water soluble is not better the water soluble is not better that pyrrolidone, hydroxypropylcellulose, polyethyleneglycol,

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PATENT ABSTRACTS OF JAPAN

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(54) PREPARATION OF SLOW-RELEASING MEDICINE

(57)Abstract:

PURPOSE: To obtain slow-releasing medicine which keeps the rate of release in the body constant irrespective of the pH in the digestive organs, by coating an easily disintegrable particle containing a principal medicine, with a layer which can easily be dissolved in body fluids, and forming the coated particles into a tablet.

CONSTITUTION: Easily disintegrable particle comprising a principal medicine and a disintegrating agent, is coated with a layer comprising wax, a water-soluble polymer and a nonionic surface-active agent having HLB of less than 9. The layer is water- soluble irrespective of the pH, and has a low water-permeability. The covered particles are formed into tablets having sufficiently high strength to endure the usual treatment. Although eluation retarding rate of each covered particle is as short as 1W10min, a slow-releasing medicine which releases the effective component slowly, at a constant rate over a long period of 7W24hr can be made by forming into tablets.

LEGAL STATUS

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